WHAT IS CLAIMED IS:

- 1. A 2,4-D monooxygenase gene in substantially pure form or a biologically active fragment thereof.
- 2. An isolated DNA sequence encoding a polypeptide having the biological activity of a 2,4-D-monooxygenase, or a nucleic acid useful as a probe specifically for and which hybridizes with such a sequence.
- 3. A DNA sequence of claim 1, having the sequence recited in Figure 10, a complementary strand thereof, a sequence differing therefrom by codon degeneracy, a sequence which hybridizes therewith and encodes a 2,4-D-monooxygenase, or a nucleic acid useful as a probe for and which hybridizes with such a sequence.
- 4. An isolated DNA sequence of claim 2, encoding a polypeptide having the biological activity of a 2,4-D-monooxygenase.
- 5. A nucleic acid sequence of claim 1 which is a 2,4-D-monooxygenase gene or a fragment thereof coding for a polypeptide having the biological activity of such a monooxygenase.
- 6. An oligonucleotide useful as a probe which hybridizes with a nucleic acid sequence of claim 1.

- 7. A oligonucleotide probe of claim 6, labelled with a detectable moiety.
- 8. A recombinant vector comprising exogenous DNA of claim 1.
- 9. A vector of claim 8, wherein the exogenous DNA is under the control of a heterologous promoter.
- 10. A substantially pure mutant menoamine exygenase gene, wherein at least one nucleic acid is substituted for, inserted into and/or deleted from a menoamine exygenase gene of claim 1, and wherein the resultant product is useful as a probe specific for the monooxygenase gene.
- 11. A substantially pure mutant monoamine exygenase, wherein at least one amino acid is substituted for, inserted into and/or deleted from a monoamine exygenase encoded by a gene of claim 1.
- 12. A mutant monoamine exygenase gene of claim 10, which is not capable of expressing a biologically active product.
- 13. A recombinant vector comprising exogenous DNA of claim 12.
- 14. A mutant plasmid comprising all genes important for degradation of 2,4-dichlorophenoxyacetic acid, wherein the gene for 2,4-D-monooxygenase has been altered such that the biological activity of 2,4-D-monooxygenase is destroyed.
 - 15. Plasmid pJP4:Tn5-2, a plasmid of claim 14.

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- 16. Plasmid pJP4:Tn5-4, a plasmid of claim 14.
- Plasmids pVJH21, pGJS3, pKJS31, pKJS32, pKJSB330, pKJS(x)630 and pKJS32RHΔS', each a vector of claim 8.
- 18. Plasmids pTRJS'B435, pTJS'B436, pTJSS'035, pTJSS'036, pTJS x535 and pTJS'x536, each a vector of claim 9.
- 19. Phages MJSS'030 and MJSS'031, each a vector of claim 13.
 - 20. Plasmid pKJE B130, a vector of claim 13.
 - 21. Plasmid pTJS'x535omega, a vector of claim 13.
- 22. A method of identifying and isolating a vector containing a 2,4-D-monooxygenase gene comprising growing on selective media a bacterium transformed with a plasmid of claim 14 and cotransformed with a vector to be tested.

A bacterial strain transformed with a vector of claim 8.

A bacterial strain transformed with a vector of claim 12

- 25. A recombinant vector comprising exogenous DNA of claim 10.
- 26. A recombinant vector comprising exogenous DNA of claim 25.
 - 27. An E. coli strain of claim 26.

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- 28 A Pseudomonas strain of claim 26.
- 29. An Alcaligenes strain of claim 26.
- 30. An Agrobacterium strain of claim 26.
- 31. Plasmids pUJC1001, pUJC01003, pmCJ1007 and pmLJC1005, each a vector of claim 22.
- 32. A transgenic microorganism or plant comprising an exogenous DNA sequence of claim 1.
- 33. A transgenic plant wherein the exogenous DNA is a vector of claim 25.
- 34. A transgenic plant wherein the exogenous DNA is a plasmid of claim 31.
- 35. A transgenic microorganism or plant comprising cells of a microorganism or plant transformed with a vector of claim 9.
 - 36. A transgenic microorganism or plant comprising cells of a microorganism or plant transformed with a vector of claim 25.
 - 37. An antibody capable of binding to 2,4-D-monooxygenase
 - 38. A monoclonal antibody of claim 37.
 - 39. A fragment of an antibody of claim 37, which fragment is capable of binding 2,4-D-monooxygenase.

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40. An antibody fragment of claim 39, which is the Fab fragment.

41. An antibody of claim 37, which is labelled.

having the biological activity of monoamine oxygenase, comprising culturing a cell of claim \$2.50

As. A method of preparing a polypeptide having the biological activity of monogmine oxygenese, comprising culturing a cell of claim 28.

44. A method of preparing a polypeptide having the biological activity of menoamine oxygenase, comprising culturing a cell of claim 26.